



The Honorable Q. Todd Dickinson
Deputy Assistant Commissioner for
Patent Policy and Projects
Office of the Assistant Commissioner for Patents
U.S. Patent and Trademark Office
Crystal Park Building 2, Suite 919
Washington, DC 20231

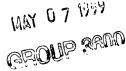
Food and Drug Administration Rockville MD 20857

RECEIVED Re: Cook GRII Coronary Stent®

RECEIVED Docket No. 97E-0461

GROUP 1700

APR 13 1000



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MAY 1 2 1999

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Dear Commissioner Dickinson:

This is in regard to the patent term extension application for U.S. Patent No. 5,041,126 filed by Cook Incorporated under 35 U.S.C. § 156. The patent claims the medical device Cook GRII Coronary Stent[®], premarket approval application PMA 910030.

In the August 7, 1998, issue of the <u>Federal Register</u> (63 Fed. Reg. 42408), the Food and Drug Administration published its determination of this product's regulatory review period, as required under 35 U.S.C. § 156(d)(2)(A). The notice provided that on or before February 3, 1999, 180 days after the publication of the determination, any interested person could file a petition with FDA under 35 U.S.C. § 156(d)(2)(B)(i) for a determination of whether the patent term extension applicant acted with due diligence during the regulatory review period.

The 180-day period for filing a due diligence petition pursuant to this notice has expired and FDA has received no such petition. Therefore, FDA considers the regulatory review period determination to be final.

Please let me know if we can provide further assistance.

Sincerely yours,

Ronald L. Wilson, Director Health Assessment Policy Staff

Office of Health Affairs

cc: C. David Emhardt

Woodward, Emhardt, Naughton, Moriarty & McNett

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